

Scientific & Regulatory Consultants, Inc.

SUBMITTED VIA CDX

November 3, 2020

Mr. Demson Fuller, PM-32
U.S. Environmental Protection Agency
Document Processing Desk (AMEND)
Office of Pesticide Programs (7504P)
Room S-4900, One Potomac Yard
2777 Crystal Drive
Arlington, VA 22202

SUBJECT: Existing Product Submission of Efficacy Data to Qualify for List N
Product: GenEon™ Mineral Electrolyte, EPA Reg. No. 91112-2

Dear Mr. Fuller:

On behalf of GenEon Technologies, LLC (GenEon), enclosed is an amendment for the subject product to add an efficacy claim against Human Coronavirus Strain 229E to qualify for List N for expedited review.

MRID 51343701 is enclosed to demonstrate the product's effectiveness against Human Coronavirus Strain 229E. The submitted study (study number A30510) supports an on-label claim against Human Coronavirus Strain 229E, and qualifies the product for List N.

Per the Agency's recent request, we have included a copy of the current accepted basic CSF for the product for reference purposes. A complete list of submitted documents can be found in the attached Transmittal.

Please contact me via email (jvenable@srcconsultants.com) or phone (260.244.6270) with questions.

Sincerely,



Jamie Venable
Agent